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37. (Amended) Nitrogen heterocyclic aromatic derivative having the following structure:

(XVII)

1239. (Amended) The pharmaceutical composition comprising a nitrogen heterocyclic aromatic derivative according to claim 37 in the form of a transdermal skin patch.

A0. (Amended) The pharmaceutical composition comprising a nitrogen heterocyclic aromatic derivative according to claim 37 for intravenous administration.

Cancel claims 47 and 48 and add the following new claims:

20 49. (New) A method of reducing the humoral and cellular immunological response of a mammal comprising administering an effective amount of a nitrogen heterocyclic aromatic derivative having the following general formula:

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where X and Y are selected from N, C and CH, provided they are different from each other, or X and Y are both nitrogen;

R is hydrogen or -COR $_8$  where R $_8$  is a saturated or non-saturated C $_1$ -C $_{10}$  aliphatic hydrocarbon, R $_1$  has the following general formula:

$$R_4$$
 (II)

where  $R_3$  is hydrogen, halogen,  $C_1$ - $C_{10}$  alkyl or  $C_1$ - $C_{10}$  alkoxyl,  $R_4$  is hydrogen,  $C_1$ - $C_{10}$  alkyl or  $C_1$ - $C_{10}$  alkoxyl, or  $R_3$  and  $R_4$  together form a methylendioxy group;

 $R_2$  has the following general structure:

$$R_6$$
 $CH_2OR_5$ 
(III)

and R<sub>5</sub> is selected from:

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where  $Z=OR_7$  where  $R_7$  is a saturated or non-saturated, linear or branched  $C_1-C_{10}$  aliphatic hydrocarbon, or  $R_7$  has the following formula:

where R,  $R_1$ , X and Y are defined as above and  $R_6$  is hydrogen, halogen, alkyl or alkoxyl  $C_1$ - $C_{10}$ , or Z is NHR<sub>8</sub> where  $R_8$  is a linear or branched  $C_1$ - $C_{20}$  alkyl chain,

provided that when X=Y=N and R is H or -CONHCH<sub>2</sub>CH<sub>3</sub>, Z is not NHR<sub>8</sub> where R<sub>8</sub> is -CH<sub>2</sub>CH<sub>3</sub>, and provided that R<sub>1</sub> and R<sub>2</sub> are not located on two adjacent atoms of the heterocyclic aromatic ring, and further provided that when X=Y=N and R<sub>5</sub> is -COZ where Z=OR<sub>7</sub>, R<sub>7</sub> is a saturated or non-saturated linear or branched C<sub>5</sub>-C<sub>20</sub> aliphatic hydrocarbon.

(Amended) A method of treating a tumor susceptible to therapy comprising administering an effective amount of a nitrogen heterocyclic aromatic derivative having the following general formula:

$$\frac{1}{\sqrt{2}} \int_{R_2}^{R_1} \frac{R}{N} R_1 \qquad (I)$$

where X and Y are selected from N, C and CH, provided they are different from each other, or X and Y are both nitrogen;

R is hydrogen or -COR<sub>8</sub> where  $R_8$  is a saturated or non-saturated  $C_1$ - $C_{10}$  aliphatic hydrocarbon,  $R_1$  has the following general formula:

$$rac{1}{\sqrt{2}}$$
  $rac{1}{\sqrt{2}}$   $rac{$ 

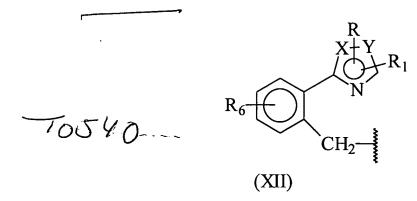
where  $R_3$  is hydrogen, halogen,  $C_1$ - $C_{10}$  alkyl or  $C_1$ - $C_{10}$  alkoxyl,  $R_4$  is hydrogen,  $C_1$ - $C_{10}$  alkyl or  $C_1$ - $C_{10}$  alkoxyl, or  $R_3$  and  $R_4$  together form a methylendioxy group;

R<sub>2</sub> has the following general structure:

$$R_6$$
 $CH_2OR_5$ 
(III)

and  $R_5$  is selected from:

where  $Z=OR_7$  where  $R_7$  is a saturated or non-saturated, linear or branched  $C_1$ - $C_{10}$  aliphatic hydrocarbon, or  $R_7$  has the following formula:



where R,  $R_1$ , X and Y are defined as above and  $R_6$  is hydrogen, halogen, alkyl or alkoxyl  $C_1$ - $C_{10}$ , or Z is NHR<sub>8</sub> where  $R_8$  is a linear or branched  $C_1$ - $C_{20}$  alkyl chain,

provided that when X=Y=N and R is H or -CONHCH<sub>2</sub>CH<sub>3</sub>, Z is not NHR<sub>8</sub> where  $R_8$  is -CH<sub>2</sub>CH<sub>3</sub>, and provided that  $R_1$  and  $R_2$  are not located on two adjacent atoms of the heterocyclic aromatic ring, and further provided that when X=Y=N and  $R_5$  is -COZ where Z=OR<sub>7</sub>,  $R_7$  is a saturated or non-saturated linear or branched  $C_5$ -C<sub>20</sub> aliphatic hydrocarbon.

51. (New) A pharmaceutical composition comprising a nitrogen heterocyclic aromatic derivative according to claim 37 together with a pharmaceutically acceptable carrier or diluent.

